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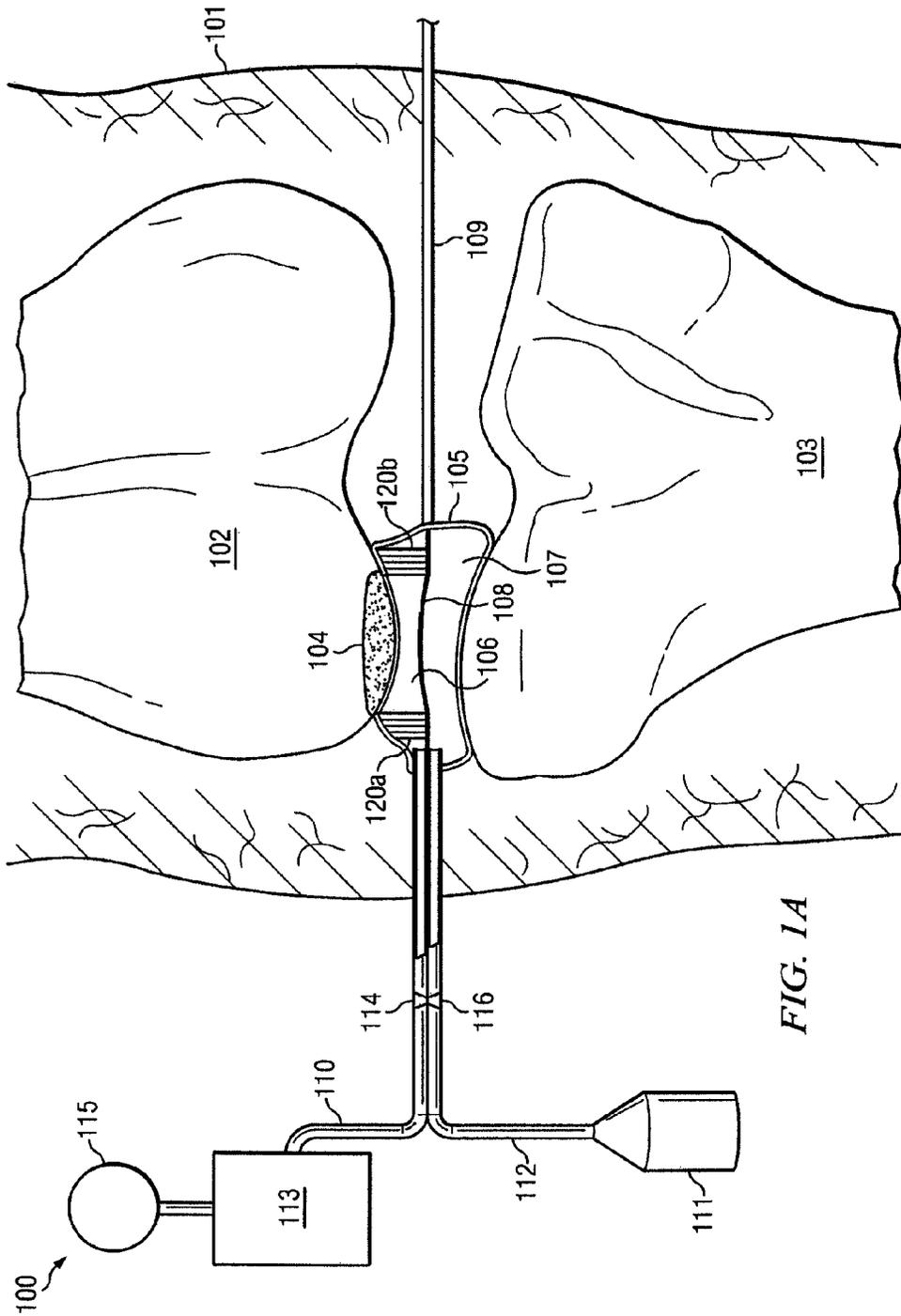


FIG. 1A

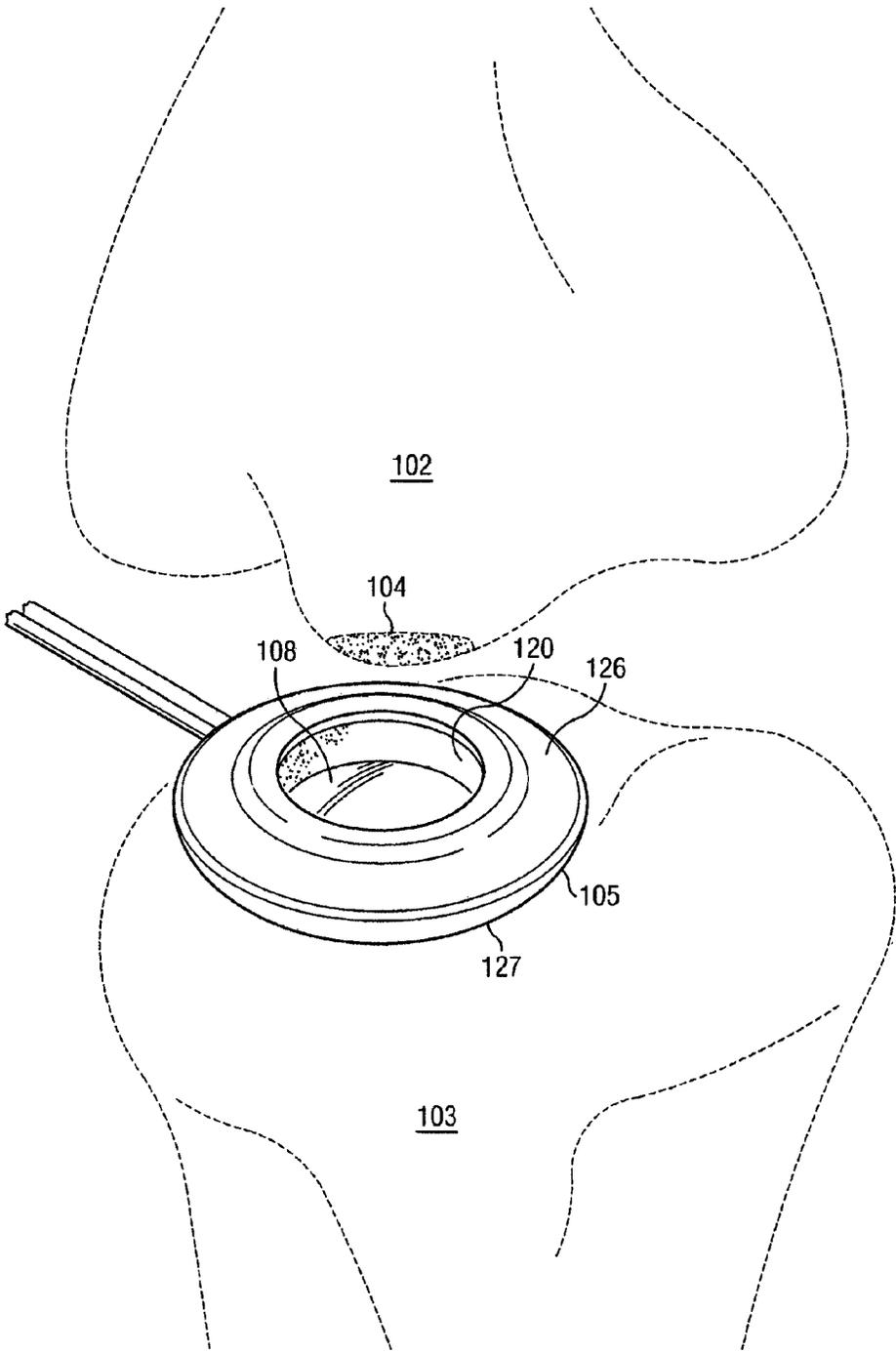
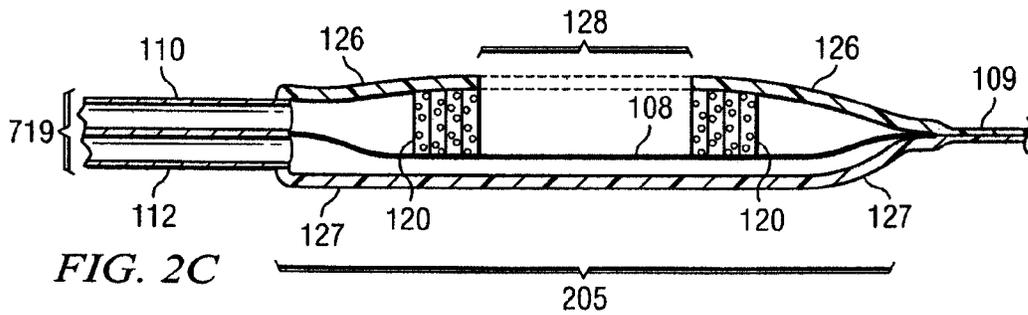
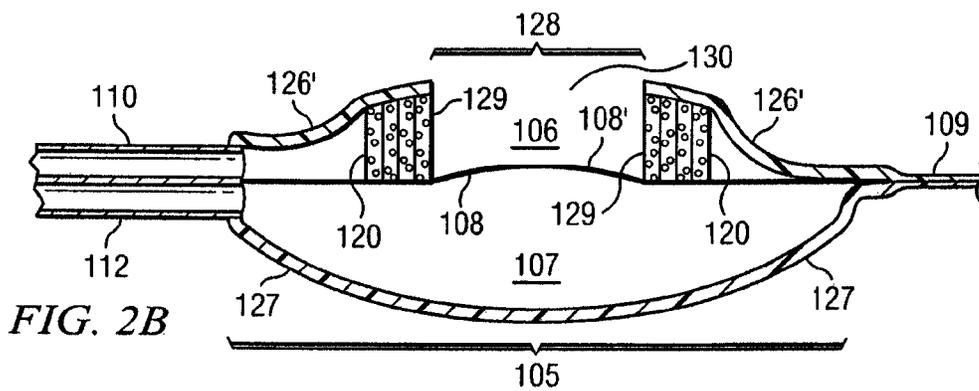
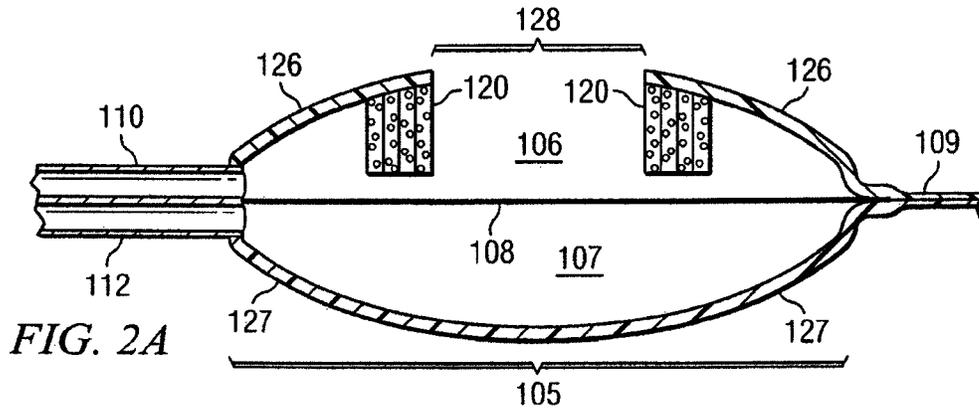


FIG. 1B



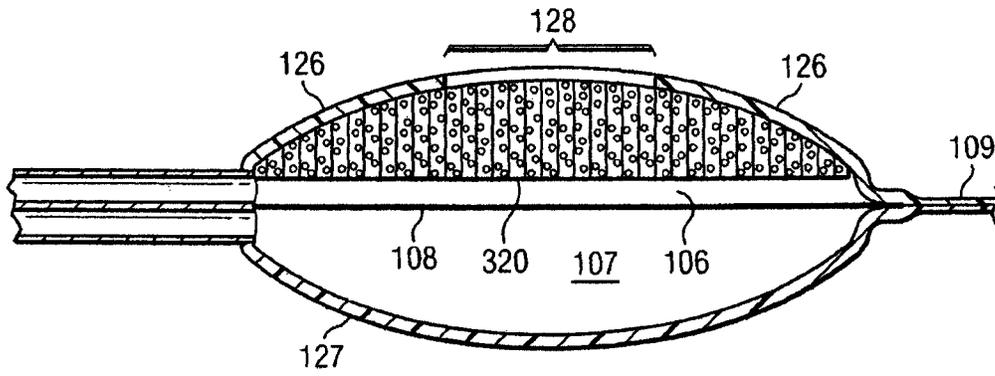


FIG. 3

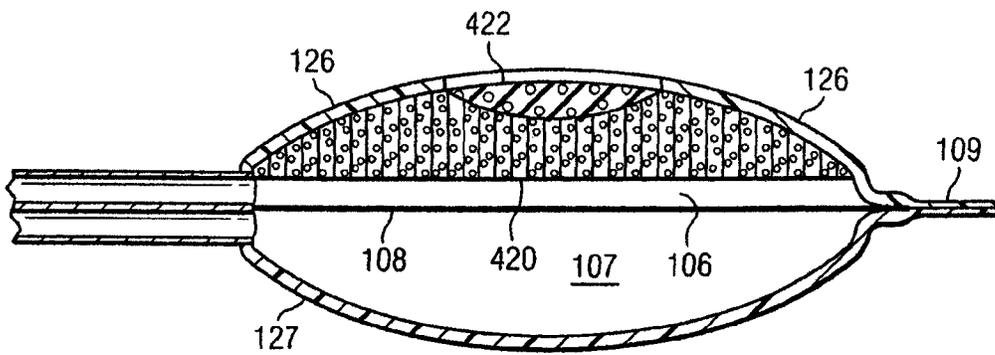


FIG. 4

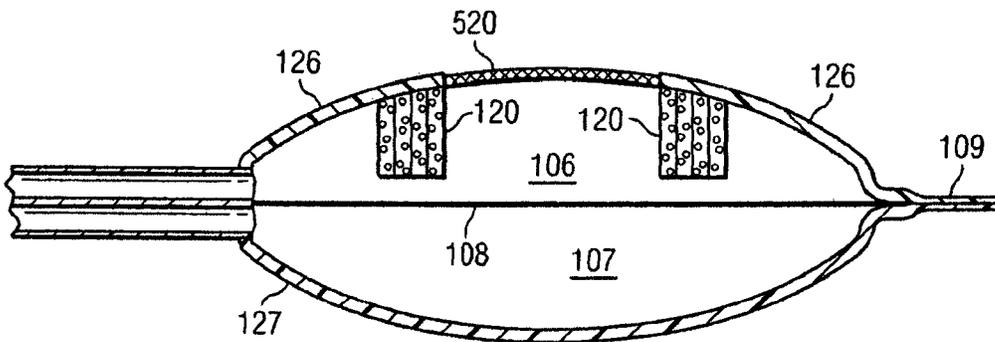


FIG. 5

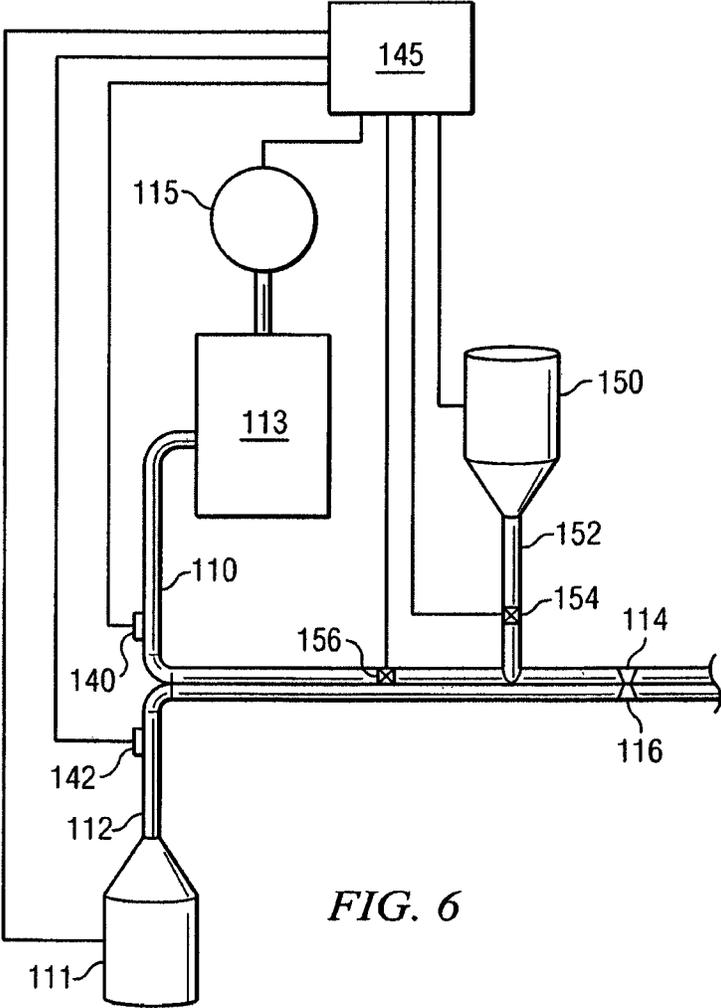
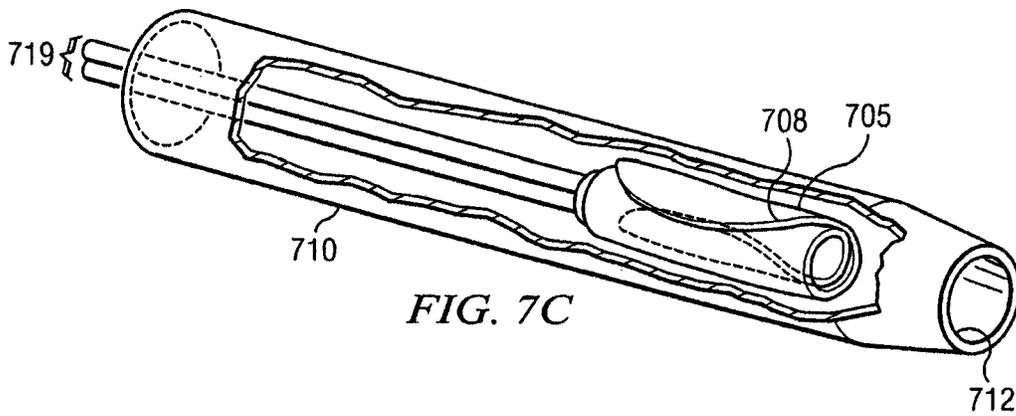
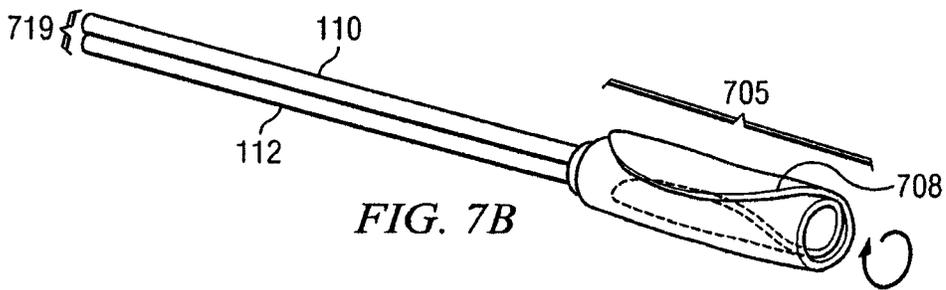
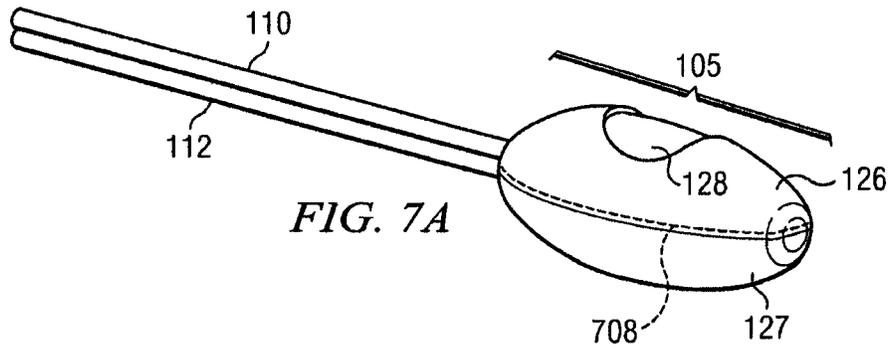


FIG. 6



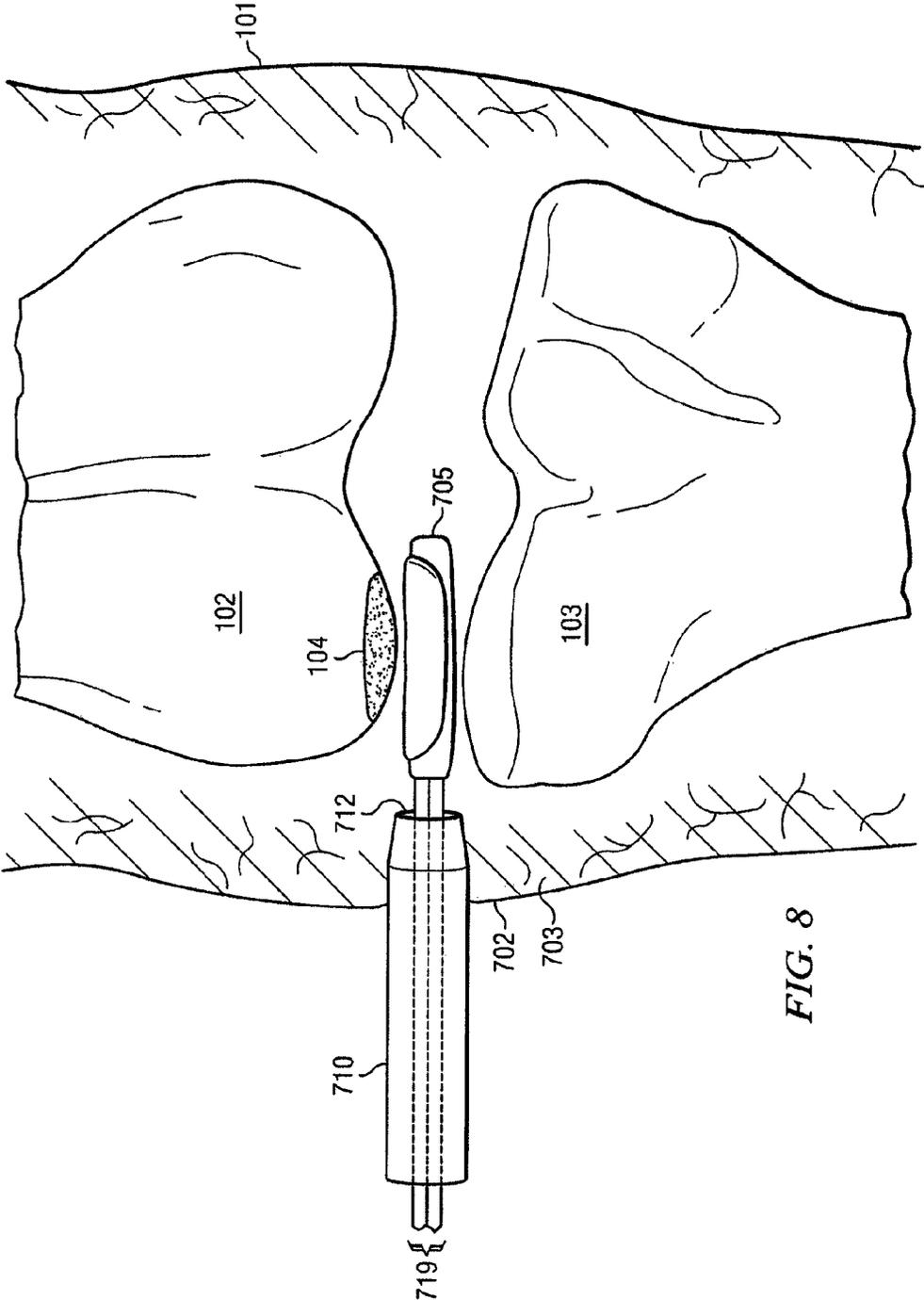


FIG. 8

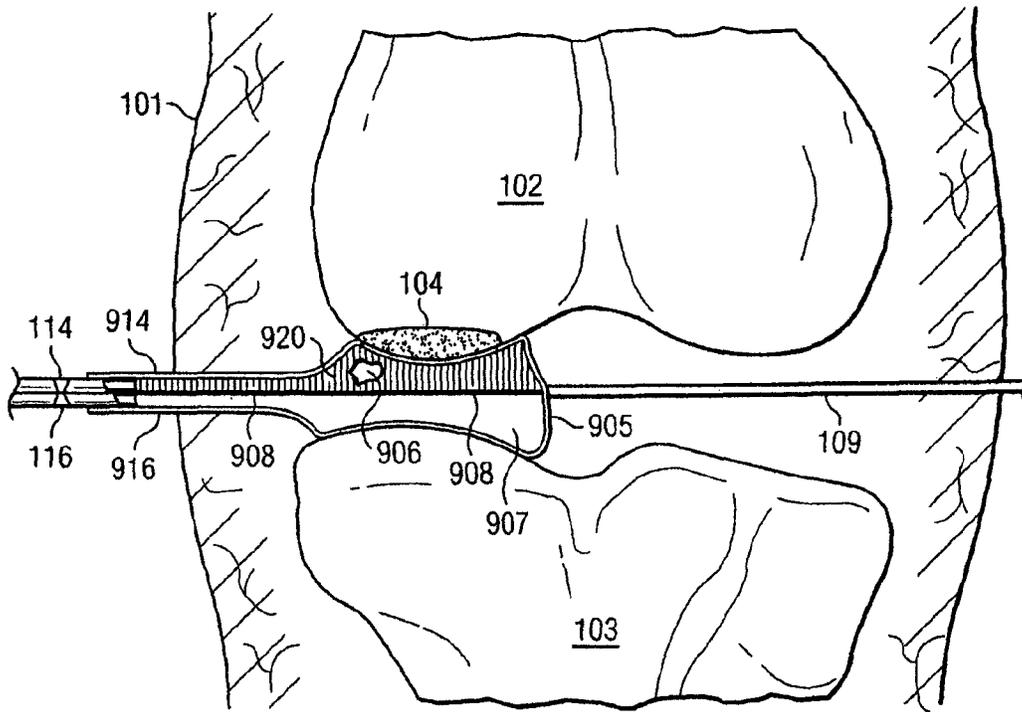


FIG. 9

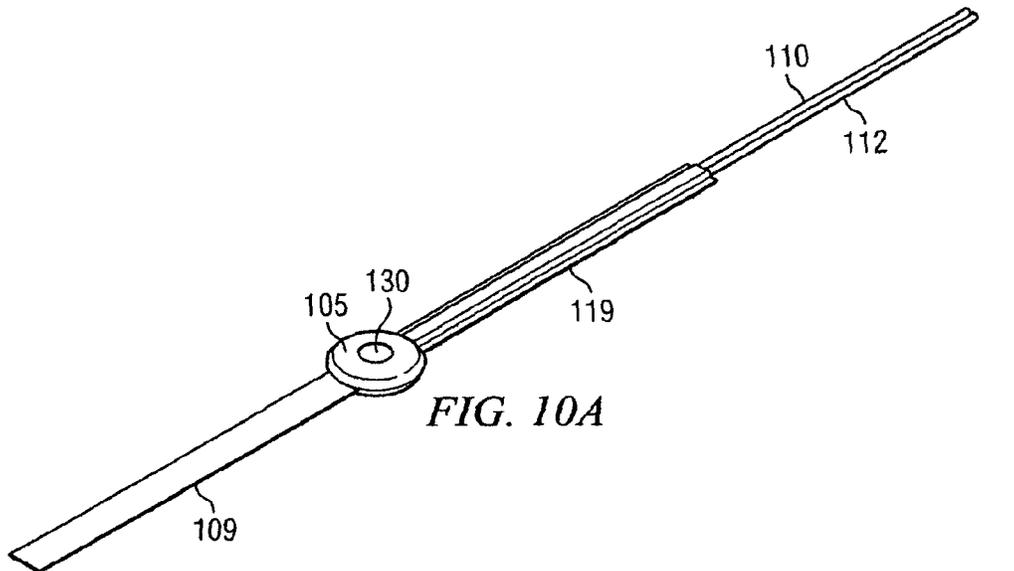


FIG. 10A

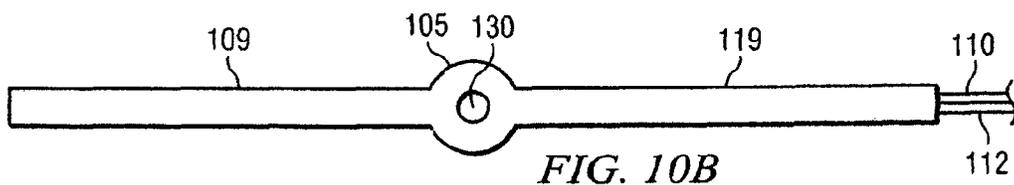


FIG. 10B

REDUCED PRESSURE AUGMENTATION OF MICROFRACTURE PROCEDURES FOR CARTILAGE REPAIR

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a divisional of U.S. application Ser. No. 12/643,508, filed Dec. 21, 2009 which claims the benefit of U.S. Provisional Application No. 61/141,593, filed Dec. 30, 2008, which is hereby incorporated by reference.

BACKGROUND

1. Field of the Invention

The present application relates generally to tissue treatment systems and in particular to treatment of articular cartilage undergoing microfracture procedure.

2. Description of Related Art

Clinical studies and practice have shown that providing a reduced pressure in proximity to a tissue site augments and accelerates the growth of new tissue at the tissue site. The applications of this phenomenon are numerous, but application of reduced pressure has been particularly successful in treating wounds. This treatment provides a number of benefits, including faster healing and increased formulation of granulation tissue. Typically, reduced pressure is applied to tissue through a porous pad or other manifolding device. The porous pad contains cells or pores that are capable of distributing reduced pressure to the tissue and channeling fluids that are drawn from the tissue. The porous pad often is incorporated into a dressing having other components that facilitate treatment.

Articular cartilage is a highly organized avascular tissue composed of chondrocytes formed in an extracellular matrix. This tissue is extremely important to the normal, healthy function and articulation of joints. Articular cartilage enables joint motion surfaces to articulate smoothly with a very low coefficient of friction. It also acts as a cushion to absorb compressive, tensile, and shearing forces and, thus, helps protect the ends of bone and surrounding tissue.

Age, injury and wear, and cartilage disorders, such as osteoarthritis, affect millions of people throughout the world. Traumatic chondral injuries, for example, are common in sports and other activities that cause severe stress and strain to joints. Osteoarthritis is also a common condition that develops as cartilage wears, weakens, and deteriorates at the joint motion surfaces of bones. Indeed, it is currently believed that 85% of all Americans will develop degenerative joint disease as a result of normal activities that damage articular cartilage.

Articular cartilage is generally thin with an extremely low or insignificant blood flow and, as such, has a very limited ability to repair or heal itself. Partial-thickness chondral defects, for example, cannot spontaneously heal. If these defects are left untreated, they often degenerate at the articular surface and progress to osteoarthritis. Full-thickness defects that penetrate subchondral bone can undergo some spontaneous repair if fibrocartilage forms at the defect. Even in spite of the formation of fibrocartilage, clinical evidence shows that full-thickness defects continue to degenerate and progress to osteoarthritis if these defects are left untreated.

Early diagnosis and treatment are crucial to hindering or stopping the progression of arthritis and degeneration of articular cartilage at joint motion surfaces. Today, depending on the grade of chondral damage, patients usually have

several surgical options to repair or regenerate articular cartilage. Some current techniques to repair cartilage include implantation of chondrocytes, implantation of synthetic matrices and surgical intervention, with reattachment and reconstruction of the damaged tissue. None of these methods are totally satisfactory and they rarely restore full function or return the tissue to its native normal state. In addition, none of these methods are proven to regenerate cartilage in situ and in vivo.

Micro-fracture surgery is one treatment modality used to treat cartilage defects. This technique is a marrow-stimulating arthroscopic procedure to penetrate the subchondral bone to induce fibrin clot formation and the migration of primitive stem cells from the bone marrow into the defective cartilage location. Generally, the base of the defective area is shaved or scraped to induce bleeding. An arthroscopic awl or pick is then used to make small holes or microfractures in the subchondral bone plate. The end of the awl is manually struck with a mallet to form the holes while care is made not to penetrate too deeply and damage the subchondral plate. The holes penetrate a vascularisation zone and stimulate the formation of a fibrin clot containing pluripotential stem cells. The clot fills the defect and matures into fibrocartilage.

While microfracture surgery has a high success rate, patients cannot return to sports or other intense activities for about 4 months, even with the help of physical therapy. Additionally, the tissue that forms in the defect is primarily fibrocartilage (this constitutes a repair process whereby a different tissue type is formed), which does not have the same functional characteristics of articular (hyaline) cartilage. As such, there is currently an acute need for a method that leads to more of a regenerative response (forms the same type of tissue that was damaged, for example, hyaline cartilage) rather than a repair response and that reduces the overall time of healing.

It would therefore be advantageous to provide devices, methods and systems to promote healing and/or tissue regeneration after microfracture surgery. Such devices, methods and systems would decrease healing time, and lead to better functional outcomes thus increasing the patient's quality of life and enable a more rapid return to normal daily activities.

SUMMARY

The problems presented by existing methods for microfracture surgery are solved by the systems and methods of the illustrative embodiments described herein. These systems and methods are designed to deliver reduced pressure to the joint space where microfracture surgery is performed. Traditional reduced pressure delivery methods to a joint space, in particular a knee joint, would apply reduced pressure to the entire joint. The systems and methods described herein concentrate application of the reduced pressure on the defect (e.g., the site of the microfracture surgery) by utilizing a bladder having a reduced-pressure chamber with an opening that surrounds the defect to avoid applying reduced pressure to the entire joint.

In one embodiment, a system for applying a treatment to a defect in a first bone of two bones forming a joint is disclosed and comprises a reduced-pressure source for providing a reduced pressure, a positive-pressure source for providing a positive pressure, and a bladder. The bladder is formed with a reduced-pressure chamber in fluid communication with the reduced-pressure source and a bracing chamber in fluid communication with the positive-pressure source. The reduced-pressure chamber and bracing chamber

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both have walls formed from a flexible material, and the reduced-pressure chamber has an opening sized to substantially surround the defect in the first bone. A portion of the walls of each of the reduced-pressure chamber and the bracing chamber form an interior wall of the bladder.

The system further comprises a manifold positioned within the reduced-pressure chamber between the interior wall and the opening in the reduced-pressure chamber and formed of a porous material for distributing reduced pressure and providing structural support between the first bone and contact portion of the interior wall. When positive pressure is applied to the bracing chamber, the walls of the bracing chamber inflate to provide bracing between the interior wall and a second bone of the two bones. When reduced pressure is applied to the reduced-pressure chamber, the walls of the reduced-pressure chamber collapse toward the interior wall causing the manifold to provide a seal with the contact portion of the interior wall against the defect and provide bracing between the contact portion of the interior wall and the first bone.

In another embodiment, a method for applying the treatment is also disclosed and first comprises performing surgery on the defect in the first bone. The method then comprises positioning a reduced-pressure chamber in the joint adjacent the defect, positioning a bracing chamber in the joint between the reduced-pressure chamber and a second bone of the two bones, and positioning a manifold within the reduced-pressure chamber adjacent the defect and the bracing chamber. The method further comprises applying a positive pressure to the bracing chamber to force the manifold against the first bone to brace the reduced-pressure chamber against a second bone of the two bones and then applying a reduced-pressure to the defect through a hole in the reduced-pressure chamber and the manifold.

In a further embodiment, a method is provided that includes performing microfracture surgery to a knee. The method comprises creating at least one microfracture in a bone at the base of an articular cartilage defect in the knee, then applying reduced pressure to the site of the defect.

Other objects, features, and advantages of the illustrative embodiments will become apparent with reference to the drawings and detailed description that follow.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1A is a schematic, cross-sectional view of a reduced-pressure treatment system for repairing cartilage in a knee joint that includes a fluidic system, a bladder positioned between the femur and tibia of the knee joint, and a first embodiment of a manifold structure within the bladder;

FIG. 1B is a schematic, perspective view of the bladder and femur shown in FIG. 1A;

FIGS. 2A and 2B are schematic, cross-sectional views of the bladder shown in FIG. 1A with and without reduced pressure being applied to the bladder;

FIG. 2C is a schematic, perspective view of the bladder shown in FIG. 1A without reduced pressure and flattened to be rolled as shown in FIG. 7B;

FIG. 3 is a schematic, cross-sectional view of the bladder shown in FIG. 1A including a second embodiment of a manifold structure;

FIG. 4 is a schematic, cross-sectional view of the bladder shown in FIG. 1A including scaffold material;

FIG. 5 is a schematic, cross-sectional view of the bladder shown in FIG. 1A including a barrier material;

FIG. 6 is a schematic view of a fluid control system for the fluidic system shown in FIG. 1A;

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FIG. 7A is a schematic, perspective view of the bladder shown in FIG. 1A, FIG. 7B is a perspective view of the bladder after being rolled or folded, and FIG. 7C is a perspective view of the rolled bladder inserted within a delivery catheter;

FIG. 8 is a schematic, cross-sectional view of the delivery catheter and rolled bladder of FIG. 7C inserted into the knee joint;

FIG. 9 is a schematic, cross-sectional side view of another embodiment of a bladder similar to the bladder shown in FIG. 1A; and

FIGS. 10A and 10B include a perspective and plan views, respectively, of the bladder shown in FIG. 1A including an elongated tab.

DETAILED DESCRIPTION OF ILLUSTRATIVE EMBODIMENTS

In the following detailed description of the illustrative embodiments, reference is made to the accompanying drawings that form a part hereof. These embodiments are described in sufficient detail to enable those skilled in the art to practice the invention, and it is understood that other embodiments may be utilized and that logical structural, mechanical, electrical, and chemical changes may be made without departing from the spirit or scope of the invention. To avoid detail not necessary to enable those skilled in the art to practice the embodiments described herein, the description may omit certain information known to those skilled in the art. The following detailed description is, therefore, not to be taken in a limiting sense, and the scope of the illustrative embodiments are defined only by the appended claims.

Referring to FIGS. 1A and 1B, a reduced pressure therapy system 100 for applying reduced pressure and facilitating the growth of tissue at a tissue site in the body of a patient such as, for example, the knee joint in a leg 101 of the patient comprising the femur 102 and tibia 103. Any portion of the knee joint may have developed a defect that needs repair, for example, a defect 104 in the articular cartilage of the lateral condyle of the femur 102. The defect 104 may also be a micro-fracture surgery site as described above where it is desired to regenerate, repair, add or promote growth of new cartilage tissue or any tissue functioning as a structure and support of the body including, without limitation, areola connective tissue, dense connective tissue and cartilage.

The reduced pressure therapy system 100 also comprises a reduced pressure source 115 for providing a reduced pressure to a bladder 105 comprising a reduced-pressure chamber 106 and a bracing chamber 107 divided by an interior wall 108 within the bladder 105. It should be understood that the chambers 106,107 may be separate chambers without the common interior wall 108. The bladder 105 is positioned between the bones 102, 103 with the reduced-pressure chamber 106 positioned adjacent the femur 102. It should be understood that the defect 104 could also be in the tibia 103 with the reduced-pressure chamber 106 positioned adjacent such defect in the tibia 103. The reduced pressure source 115 is fluidly coupled to the reduced-pressure chamber 106 of the bladder 105 via a first conduit 110. The reduced pressure therapy system 100 further comprises a canister 113 fluidly coupled between the reduced pressure source 115 and the reduced pressure chamber 106 of the bladder 105 to collect bodily fluids, such as blood or exudate, that are drawn from the femur 102 via the

defect **104**. In one embodiment the reduced-pressure source **115** and the canister **113** are integrated into a single housing structure.

In the context of this specification, the term “reduced pressure” generally refers to a pressure that is less than the ambient pressure at a tissue site that is subjected to treatment. In most cases, this reduced pressure will be less than the atmospheric pressure of the location at which the patient is located. Although the terms “vacuum” and “negative pressure” may be used to describe the pressure applied to the tissue site, the actual pressure applied to the tissue site may be significantly greater than the pressure normally associated with a complete vacuum. Consistent with this nomenclature, an increase in reduced pressure or vacuum pressure refers to a relative reduction of absolute pressure, while a decrease in reduced pressure or vacuum pressure refers to a relative increase of absolute pressure.

The system **100** further comprises a first fluid supply **111** that is fluidly coupled to the bracing chamber **107** of the bladder **105** via a second conduit **112**. The first fluid supply **111** provides a bracing substance to the bracing chamber **107** via the second conduit **112** to fill the bracing chamber **107** under a positive pressure so that the bracing chamber **107** expands and exerts a positive force on the tibia **103** and surrounding tissue as well as the interior wall **108** of the bladder **105**. The positive pressure applied to the bracing substance in the bracing chamber **107** is sufficient to allow the bracing chamber **107** to conform to the shape and contours of the tibia **103** and surrounding tissue while simultaneously providing a cushion between the femur **102** and the tibia **103** without collapsing. It is to be understood that the bracing chamber **107** of the bladder **105** may comprise multiple chambers to achieve the desired cushioning and conformance with the tibia **103** and surrounding tissue. The bracing substance may be a gas or a liquid such as, for example, a highly viscous compressible material such as a putty, slurry, or colloid.

Both the first conduit **110** and the second conduit **112** may be coupled to the bladder **105** via connectors **114** and **116**, respectively. Both conduits **110**, **112** may be coated with an anti-coagulant to prevent a build-up of bodily fluids or blood within the conduits. As used herein, the term “coupled” includes direct coupling or indirect coupling via a separate object. The term “coupled” also encompasses two or more components that are continuous with one another by virtue of each of the components formed from the same piece of material. Also, the term “coupled” may include chemical, mechanical, thermal, or electrical coupling. Fluid coupling means that fluid is in communication between the designated parts or locations.

The bladder **105** is shown with and without a reduced pressure applied to the reduced-pressure chamber **106** in FIGS. 2B and 2A, respectively. Referring more specifically to FIG. 2A, the bladder **105** comprises two exterior walls including a reduced-pressure exterior wall **126** which forms the external surface of the reduced-pressure chamber **106** and bracing exterior wall **127** which forms the external wall of the bracing chamber **107**. The reduced-pressure exterior wall **126** has a generally circular opening **128** which is shaped to fit the perimeter of the defect **104**. The bladder **105** further comprises a manifold structure **120** that may be two separate manifold components **120a**, **120b** positioned on both sides of the defect **104** in the femur **102** as shown in FIG. 1A or that may be substantially cylindrical in shape with an inner diameter that is concentrically aligned with the opening **128** of the reduced-pressure exterior wall **126** as shown in FIG. 1B. The upper surface of the manifold

structure **120** may be sealed against the inside surface of the reduced-pressure exterior wall **126**. The manifold structure **120** may have a height somewhat less than the distance between the reduced-pressure exterior wall **126** and the interior wall **108** so that it is slightly suspended over the interior wall **108** when the reduced-pressure chamber **106** is not under pressure.

When a reduced pressure is applied to the reduced-pressure chamber **106** as shown in FIG. 2B, the reduced-pressure exterior wall **126** collapses to the position shown as **126'** with the manifold structure **120** being forced against the interior wall **108** and forming a seal with a portion of the interior wall **108** to concentrate the reduced pressure on the defect **104**. The portion of the interior wall **108** defined by the inner diameter of the manifold structure **120** may form a slightly dimpled portion **108'** as a result of the reduced pressure being applied through the manifold structure **120**. The dimpled portion **108'** of the interior wall **108** and the inner surface **129** of the manifold structure **120** define a cavity **130** that is positioned adjacent the defect **104**. Consequently, the reduced-pressure exterior wall **126** provides a seal around the defect **104** for applying the reduced pressure directly to the defect **104** via the opening **128** in the reduced-pressure exterior wall **126**. The manifold structure **120** also serves to prevent the exterior wall **126** from collapsing against the interior wall **108**, allowing the reduced pressure to continue to be applied to the defect **104** in the femur **102**. The manifold structure **120**, whether a single piece or multiple manifold components **120a** and **120b**, is selected to provide adequate structural support within the reduced-pressure chamber **106** between the femur **102** and the interior wall **108** while still functioning within the reduced-pressure chamber **106** as forming a part of the cavity **130**.

The interior wall **108** is substantially impermeable to block the transmission of fluids including both liquids and gases. The interior wall **108** may be a single layer of material or multiple layers of material depending on the manufacturing process of the bladder **105**. Although the manifold structure **120** may form a slight dimpled portion **108'** as a result of the reduced pressure being applied to the reduced-pressure chamber **106** and the positive pressure being applied to the bracing chamber **107**, the interior wall **108** is substantially inflexible to reduce the amount of bending or flexing in response to the pressures being applied on both sides. Such materials may include, for example, fiberglass, metals, rubbers, or plastics alone or in combination. The interior wall **108** must be sufficiently inflexible or rigid to ensure that the cavity **130** does not collapse against the defect **104** when the reduced pressure and bracing pressure are applied to the bladder **105**.

The exterior walls **126**, **127** are also fabricated from an impermeable material to substantially block the transmission of fluids. Additionally, the exterior walls **126**, **127** must be sufficiently thick to withstand the tensile stress and compressive stress created by the application of the reduced pressure to the reduced-pressure chamber **106** and the positive pressure to the bracing chamber **107**, respectively. Consequently, the exterior walls **126**, **127** may be formed of multiple layers of material having the same or different properties. For example, the exterior walls **126**, **127** may have a first layer that is impermeable to fluids and second layer that has sufficient flexibility and structural support when under pressure. These characteristics are of course dependent upon the specific materials used to fabricate the exterior walls **126**, **127** of the bladder **105**.

The material used to form the exterior walls **126**, **127** may contain one or more elastomers so that they have rubber-like properties. For example, in some embodiments, the flexible material has elongation rates greater than 100% and a significant amount of resilience. The resilience of a material refers to the material's ability to recover from an elastic deformation. Examples of elastomers include, but are not limited to, natural rubbers, polyisoprene, styrene butadiene rubber, chloroprene rubber, polybutadiene, nitrile rubber, butyl rubber, ethylene propylene rubber, ethylene propylene diene monomer, chlorosulfonated polyethylene, polysulfide rubber, polyurethane, and silicones. Elastomers may also include, but are not limited to, polyurethane elastomers, including elastomers based on both aromatic and aliphatic isocyanates; flexible polyolefins, including flexible polyethylene and polypropylene homopolymers and copolymers; styrenic thermoplastic elastomers; polyamide elastomers; polyamide-ether elastomers; ester-ether or ester-ester elastomers; flexible ionomers; thermoplastic vulcanizates; flexible poly(vinyl chloride) homopolymers and copolymers; flexible acrylic polymers; and blends and alloys of these, such as poly(vinyl chloride) alloys like poly(vinyl chloride)-polyurethane alloys. The exterior walls **126**, **127** may also include, but are not limited to, polyester-polyurethanes, polyether-polyurethanes, and polycarbonate-polyurethanes. The different elastomeric materials described above may be combined as blends to form the exterior walls **126**, **127** in one layer, or alternatively, may be formed as separate layers. The exterior walls **126**, **127** may be formed of bioinert materials, i.e., materials which do not elicit a strong immunological reaction against the material and are not toxic, but which do not degrade within the body over time.

The manifold structure **120** distributes the reduced pressure within the reduced-pressure chamber **106** to the defect **104** while providing support as a spacer between the interior wall **108** and the reduced-pressure exterior wall **126** in the collapsed position **126'** as shown in FIG. 2B. When the reduced-pressure exterior wall **126** is forced against the femur **102**, the reduced pressure is applied directly to the defect **104** via the cavity **130** which includes the manifold structure **120** that continues to distribute the reduced pressure to the defect **104**. The manifold structure **120** comprises an open-cell foam material that includes a plurality of cells fluidly coupled to each other to form a plurality of flow channels within the manifold structure **120**. The cells and flow channels may be of uniform shape and size or may include a pattern or random variations to more precisely direct the flow of fluids through the flow channels within the manifold structure **120**.

In one illustrative embodiment, the manifold structure **120** is a foam material that may be either hydrophobic or hydrophilic. In one non-limiting example, the manifold structure **120** is an open-cell, reticulated polyurethane foam such as GranuFoam® dressing available from Kinetic Concepts, Inc. of San Antonio, Tex. In the example in which the manifold structure **120** is made from a hydrophilic material, the manifold structure **120** also functions to wick fluid out of the cavity **130**, while continuing to provide reduced pressure to the defect **104** as a manifold. The wicking properties of the manifold structure **120** draw fluid away from the defect **104** by capillary flow or other wicking mechanisms. An example of a hydrophilic foam is a polyvinyl alcohol, open-cell foam such as V.A.C. WhiteFoam® dressing available from Kinetic Concepts, Inc. of San Antonio, Tex. Other hydrophilic foams may include those made from polyether.

Other foams that may exhibit hydrophilic characteristics include hydrophobic foams that have been treated or coated to provide hydrophilicity.

In another embodiment, the manifold structure **120** may be constructed from bioresorbable materials that do not have to be removed from a patient's body after the defect **104** has been repaired and the reduced-pressure chamber **106** fully collapses. Suitable bioresorbable materials may include, without limitation, a polymeric blend of polylactic acid (PLA) and polyglycolic acid (PGA). The polymeric blend may also include, without limitation, polycarbonates, polyfumarates, and capralactones. The manifold structure **120** may further serve as a scaffold for new cell-growth, or a scaffold material may be used in conjunction with the manifold structure **120** to promote cell-growth. A scaffold is a substance or structure used to enhance or promote the growth of cells or formation of tissue, such as a three-dimensional porous structure that provides a template for cell growth. Illustrative examples of scaffold materials include synthetic, biosynthetic, and biologic material such as calcium phosphate, collagen, PLA/PGA, coral hydroxy apatites, carbonates, autologous tissue, or processed allograft or xenograft materials.

The manifold structure **120** may also include a closed-cell material to provide sufficient support within the reduced-pressure chamber **106** to prevent the exterior wall **126** collapsing prematurely on the interior wall **108**. The closed-cell material also contains a plurality of cells, but the majority of these cells are not fluidly coupled to each other with flow channels so that they provide a more solid structural support within the reduced-pressure chamber **106**. The closed-cell material is sufficiently dense to maintain the three-dimensional shape of the manifold structure **120** to provide structural support within the reduced-pressure chamber **106** and to prevent the open-cells from collapsing and cutting off the flow of reduced pressure through the flow channels.

Consequently, the size of the cells in the manifold structure **120** must be sufficiently large to facilitate the communication of fluids through the plurality of cells in the manifold structure **120**, but sufficiently small with lower porosities to provide structural support within the reduced-pressure chamber **106**. While the size of the cells in the manifold structure **120** is not exactly known, it is between about 100 microns at the low end to maintain sufficient permeability allowing air and fluids to move therethrough when under pressure, and about 1500 microns on the high end to maintain sufficient structural support for the femur **102**. Further, the size and number of cells in the manifold structure **120** affect the porosity of the manifold structure **120**. In one embodiment, the porosity percentage should be at least about 50% to ensure that the manifold structure **120** continues to facilitate fluid flow through the open-cell structure. The porosity percentage should be no greater than about 85% to ensure that the manifold structure provides sufficient structural support. Thus, the porosity of the manifold structure **120** should be in a range of about 50% to about 85%.

Referring again to FIGS. 1A, 1B, and 2B, the interior wall **108** is sufficiently rigid to prevent the dimpled portion **108'** of the interior wall **108** and the inner diameter surface **129** of the manifold structure **120** from contacting the defect **104**. Material may be positioned within the cavity **130** adjacent the defect **104** that promotes healing of the defect **104**, or provides additional structural support, or provides a barrier to isolate the reduced-pressure chamber **106** from the defect **104**, or all the foregoing. For example, additional manifold

material may be placed within the cavity **130** to provide additional structural support in a similar fashion as the manifold structure **120** described above, and/or to function as scaffold material to promote healing of the defect **104**. A manifold structure may also function as a scaffold material to promote healing of the defect **104**. Such a manifold structure may be bioresorbable and left in place after the reduced pressure delivery system **100** is removed.

The shape of the manifold structure **120** may be any that prevents the exterior wall **126** from collapsing against the interior wall **108** within the reduced-pressure chamber **106** when being subjected to a reduced pressure. Thus, the manifold structure **120** may support only a portion of the exterior wall **126** of the reduced-pressure chamber **106** as illustrated in FIG. 2A. Alternatively, a manifold structure **320** may fill most of the reduced-pressure chamber **106** to support the entire inner portion of the exterior wall **126** of the reduced-pressure chamber **106** as shown in FIG. 3. When a reduced pressure is applied to the reduced-pressure chamber **106**, the reduced-pressure exterior wall **126** does not collapse to the position shown as **126'** as shown in FIG. 2B. However, the manifold structure **320** is still forced against the interior wall **108** to form the cavity **130** positioned adjacent the defect **104**. Consequently, the reduced-pressure exterior wall **126** provides a seal around the defect **104** for applying the reduced pressure directly to the defect **104** via the opening **128** in the reduced-pressure exterior wall **126** as shown in FIG. 2B. This particular embodiment provides more structural support to the reduced-pressure chamber **106** of the bladder **105** if, for example, more support is needed for the femur **102** as preferred in some embodiments.

The manifold structure **320** may also include a portion that functions as scaffold material to promote healing of the defect **104**. The scaffold portion may be sized to fill a defect that is a gap or void in the femur **102** and may be a separate or integrated component of the manifold structure **320**. The scaffold portion of the manifold structure **320** may be bioresorbable and left in place after reduced-pressure delivery system **10** is removed. Referring more specifically to FIG. 4, such a manifold structure **420** is shown that includes a scaffold component **422** to promote regeneration of the defect **104** while at the same time providing additional structural support between the femur **102** and the interior wall **108** of the bladder **105**. The manifold structure **420** may be constructed of similar foam material as the manifold structure **120** as described above. The scaffold component **422** comprises primarily open-cell material to facilitate fluid communication between the manifold structure **420** and the defect **104**. As such, the scaffold component **422** serves as an additional manifold to more efficiently distribute reduced pressure across the entire surface of the defect **104**, while at the same time providing some cushion between the femur **102** and the interior wall **108** of the bladder **105**. The scaffold component **422** of the manifold structure **420** may be coated or infused with growth factors, cells or nutrients to promote the growth of cells and/or the regeneration of cartilage tissue. The scaffold component **422** may also be self-seeded whereby the reduced pressure pulls cells from the surrounding bone or other body tissues into the cells of the scaffold component **422** where they may grow and/or form new tissue. Further, the scaffold component **422** may be seeded with cells *ex vivo*, prior to application.

The size of the cells in the scaffold component **422** must be sufficiently large to maintain fluid flow through the plurality of cells in the structure, but sufficiently small with lower porosities to provide enough structural support within the scaffold component **422** to ensure that the flow channels

therein do not collapse significantly and the scaffold component **422** retains its shape to function as a scaffold when a reduced pressure is applied to the reduced-pressure chamber **106**. The scaffold component **422** in a reduced pressure environment may have the open cell structure that collapses to about 10%, about 20%, about 40%, about 80%, or any range therebetween to avoid disrupting or significantly altering fluid communication within the scaffold component **422**. The cell size and porosity of the scaffold component would be in the same range as described above for the manifold structure **120**.

The scaffold component **422** may be constructed as a separate component that contacts a surface of the manifold structure **120** with a loose interface that allows the scaffold component **422** to separate easily from the manifold structure **120** when the bladder **105** is removed from the femur **102**. Thus, the scaffold component **422** remains in place at the site of the repaired defect **104** when the bladder **105** is removed. Alternatively, the scaffold component **422** may be constructed so that it adheres to the surface of the manifold structure **120** using, for example, an adhesive that causes the scaffold component **422** to remain attached to the manifold structure **120** when the bladder **105** removed from the femur **102**. Thus, the scaffold component **422** does not remain in place at the site of the repaired defect **104** when the bladder **105** is removed. When the scaffold component **422** is intended to be removed with the manifold structure **420**, they may be fabricated as a unitary component (not shown) with dual porosities such that they function in the same manner as described above. Such unitary components with dual porosities are disclosed and described in U.S. Pat. No. 6,695,823 and International Application No. PCT/US2008/000596 (International Publication No. WO 2008/091521 A2), both of which are hereby incorporated by reference herein.

The scaffold component **422** may be constructed from materials including synthetics, biosynthetics, or biologics such as calcium phosphate, collagen, PLA/PGA, coral hydroxy apatites, carbonates, autologous tissue, or processed allograft or xenograft materials. The scaffold component **422** may also be formed of bioinert materials, i.e., materials which do not elicit a strong immunological reaction against the material nor are toxic, but which do not degrade within the body over time. In another embodiment, the scaffold component **422** is formed of biocompatible materials, i.e., materials which do not elicit a strong immunological reaction against the material nor are toxic, and which degrade into non-toxic, non-immunogenic chemical species which are removed from the body by excretion or metabolism. In yet another embodiment, the scaffold component **422** is formed of biodegradable materials, i.e., materials which are enzymatically or chemically degraded *in vivo* into simpler chemical species.

In some embodiments, the scaffold component **422** is made of biocompatible and/or biodegradable materials. Thus, the scaffold component **422** may remain at the site of the defect **104** even if the bladder **105** is removed from the knee joint. When the scaffold component **422** remains in the knee joint, it should be constructed of materials having appropriate dimensions for the site and sufficient strength to support new tissue growth. The biocompatible and/or biodegradable materials may include, but is not limited to, lactide, poly(lactide) (PLA), glycolide polymers, poly(glycolic acid) (PGA), poly(lactide-co-glycolide) (PLGA), ethylene glycol/lactide copolymers, polycaprolactone, poly(p-dioxanone), polyhydroxybutyrate, polyurethanes, polyphosphazenes, poly(ethylene glycol)-poly(lactide-co-gly-

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colide) co-polymer, polyhydroxyacids, polycarbonates, polyamides, polyanhydrides, polyamino acids, polyortho esters, polyacetals, degradable polycyanoacrylates, polycarbonates, polyfumarates, degradable polyurethanes, proteins such as albumin, collagen, fibrin, synthetic and natural

polyamino acids, polysaccharides such as alginate, heparin, other naturally occurring biodegradable polymers of sugar units, processed cadaveric or non-human tissues including but not limited to dermis, pericardium, tendon, or cartilage. If the scaffold component 422 is made of biocompatible and/or biodegradable materials, the materials may be designed to degrade within a desired time frame. In one embodiment, the desired degradation time frame is one to two weeks. In another embodiment, the desired degradation time frame is between one month and one year. In yet another embodiment, the desired degradation time is greater than a year. Further, in some embodiments, scaffold component 422 made of biocompatible or biodegradable materials may degrade in a manner related to the molecular weights of the materials used to make the scaffold component 422. Thus, a higher molecular weight material may result in scaffold component 422 that retains its structural integrity for longer periods of time, while lower molecular weights result in faster degradation with shorter scaffold life.

As indicated above, the cavity 130 may include a material positioned within the void adjacent the defect 104 that promotes healing of the defect 104, provides additional structural support, and provides a barrier to isolate the reduced-pressure chamber 106 from the defect 104. FIG. 5 illustrates another embodiment of the reduced-pressure chamber 106 comprising a barrier material 520 that covers the opening in the exterior wall 126 and closes the cavity 130. The barrier material 520 may be attached to the manifold structure 120 and/or attached to the exterior wall 126 of the reduced-pressure chamber 106. Thus, the barrier material 520 directly contacts the surface of the defect 104 when the cavity 130 is placed adjacent the defect 104.

The barrier material 520 may comprise one or more pieces of material that covers a portion, or all, of the exterior wall 126 of the reduced-pressure chamber 106. The barrier material 520 is preferably fabricated from a material that allows the reduced-pressure chamber 106 to be in fluid communication with the defect 104 while isolating the defect 104 from direct communication with the cavity 130. As such, the barrier material 520 may be, for example, a mesh, a sieve, a screen, or a solid sheet with spaces or a punched pattern. The barrier material 520 may have sufficient strength to provide additional structural support for the knee joint.

Referring to FIG. 6, the reduced pressure therapy system 100 may further comprise pressure sensors 140, 142 operably connected to the first and second conduits 110, 112 to measure the reduced pressure and the bracing pressure, respectively. The system further includes a control unit 145 electrically connected to the pressure sensors 140, 142, the reduced pressure source 115, and the first fluid supply 111 that provides the bracing fluids through the second conduit 112 to the bracing chamber 107. The pressure sensor 140 measures the reduced pressure of the reduced-pressure chamber 106, and also may indicate whether the first conduit 110 is occluded with blood or other bodily fluids. The pressure sensor 140 also provides feedback to control unit 145 which regulates the reduced-pressure therapy being applied by the reduced-pressure source 115 through the first conduit 110 to the reduced-pressure chamber 106. Correspondingly, the pressure sensor 142 measures the positive pressure of the bracing fluid being applied by the first fluid

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supply 111 through the second conduit 112 to the bracing chamber 107. The pressure sensor 142 also provides feedback to the control unit 145 which regulates positive pressure therapy being applied by the reduced-pressure source 115 to the bracing chamber 107. The control unit 145 also balances the relative amount of reduced pressure and positive pressure being applied to the reduced-pressure chamber 106 and the bracing chamber 107, respectively, so that sufficient pressures are being applied to both the femur 102 and the tibia 103, respectively, and the interior wall 108 as described above.

The reduced-pressure therapy system 100 may also comprise a second fluid supply 150 fluidly coupled to the first conduit 110 via third conduit 152 and operatively connected to the control unit 145. The second fluid supply 150 may be used to deliver growth and/or healing agents to the defect 104 including, without limitation, an antibacterial agent, an antiviral agent, a cell-growth promotion agent, an irrigation fluid, or other chemically active agents. The system 100 further comprises a first valve 154 positioned in the third conduit 152 to control the flow of fluid therethrough, and a second valve 156 positioned in the first conduit 110 between the reduced-pressure source 115 and the juncture between the first conduit 110 and the third conduit 152 to control the flow of reduced pressure. The control unit 145 is electrically connected to the second fluid supply 150 and the first and second valves 154, 156 to control the delivery of reduced pressure and/or fluid from the second fluid supply 150 to the reduced-pressure chamber 106 as required by the particular therapy being administered to the patient. The second fluid supply 150 may deliver the liquids as indicated above, but may also deliver air to the reduced-pressure chamber 106 to promote healing and facilitate drainage at the site of the defect 104.

The independent paths of fluid communication provided by the conduits 110, 112, and 152 may be accomplished in a number of different ways, including that of providing a single, multi-lumen tube with two or more lumens. A person of ordinary skill in the art will recognize that the pressure sensors 140, 142, valves 154, 156, and other components associated with the conduits could also be similarly associated with a particular lumen in the delivery tube if a multi-lumen tube is used. Further, additional lumens may be provided to separately introduce air or other fluids to the site of the defect 104, including, without limitation, an antibacterial agent, an antiviral agent, a cell-growth promotion agent, an irrigation fluid, or other chemically active agents.

Referring to FIGS. 7A and 2C, the exterior walls 126, 127 of the bladder 105 are flexible as described above. Thus, the bladder 105 may be collapsed or flattened as shown more specifically in FIG. 2C as a flattened bladder 205 and preferably flattened at the edge where the interior wall 108 intersects the exterior walls 126, 127, the edges of which are indicated by the reference number 708 shown by the dashed line in FIG. 7A. The flattened bladder 205 is then folded or rolled as shown in FIG. 7B into a substantially tubular shape as indicated by reference numeral 705. It should be understood again as described above that the reduced-pressure chamber 106 and the bracing chamber 107 (not shown) may not share a common wall such as the interior wall 108, but may have separate walls. The rolled bladder 705 along with the first and second conduits 110, 112 may then be inserted into a delivery catheter 710 as shown in FIG. 7C. The delivery catheter 710 has a distal end 712 that tapers so that the delivery catheter 710 may be surgically or percutaneously inserted in the knee joint between the femur 102 and the tibia 103 to deliver the rolled bladder 705 to the desired

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lateral site adjacent the defect **104** as shown in FIG. **8**. When percutaneously inserted, the delivery catheter **710** may be inserted through a sterile insertion sheath (not shown) that penetrates the skin tissue of the patient. The distal end **712** of the delivery catheter **710** is tapered to assist movement of the delivery catheter **710** through the tissues of the knee joint, but must also have a sufficiently large opening in the tapered distal end **712** to allow the rolled bladder **705** to be pushed out of the delivery catheter **710** and inserted into the knee joint of the patient.

The delivery catheter **710** may be made of any material that provides sufficient strength to allow the delivery catheter **710** to be pushed through the bodily tissues into the knee joint of the patient. Preferably, the materials of the delivery catheter **710** should be bioinert, therefore not causing injury, toxic or immunologic reaction to the surrounding body tissues. Materials of interest includes, but are not limited to, polyimide, polyamide, PBAX™, polyethylene, fluoropolymer, polyurethane, polyisoprene, nylon, steel and the like. It is also contemplated that in one alternate embodiment, the materials of the delivery catheter **710** will allow the delivery catheter **710** an amount of flex. Further, in one embodiment, outer diameter of the delivery catheter **710** is coated with agents to prevent the tube from adhering to the body tissues. For example, the tubing may be coated with heparin, anticoagulants, anti-fibrogens, anti-adherents, anti-thrombinogens or hydrophilic substances. In another embodiment, the outer diameter and/or interior diameter of the delivery catheter **710** are coated with a lubricant agent, including but not limited to, silicone, hydrophilic coating, Surmodics® lubricants, and the like.

Referring again to FIG. **8**, the distal end **712** of the delivery catheter **710** is pushed through the skin **702** of the leg **101** of the patient until it is positioned adjacent the knee joint. The first and second conduits **110** and **112**, collectively referred to as the conduits **719**, may then be manually pushed to force the rolled bladder **705** through the delivery catheter **710** and out of the distal end **712** of the delivery catheter **710** until the rolled bladder **705** is laterally positioned at the desired site adjacent the defect **104**. As described above, the femur **102** with the defect **104** may have already undergone microfracture surgery. Although it is understood that exact equipment and method of microfracture surgery may change over time, one of skill in the art readily understands the procedure of microfracture surgery generally includes the medical practitioner, i.e., surgeon or nurse practitioner, who cuts a small incision on the skin **702** exterior to the knee joint. A long thin scope (not shown), for example, an arthroscope, may be inserted through the skin **702** and through the body tissues **703** to the site of the defect **104**. If necessary, calcified cartilage is removed. The surgeon may then create microfractures, i.e., small holes, scrapes, tears, in the femur **102** near the defective cartilage, i.e., the defect **104**. After the long thin scope is removed, the delivery catheter **710** may be inserted through the same skin opening **702** and body tissue **703** path formed by the scope. In another embodiment, a separate, additional skin opening may be formed for receiving the delivery catheter **710**. After the distal end **712** of the delivery catheter **710** is delivered to the surgical site, the conduits **719** may be mechanically or manually pushed so that the rolled bladder **705** moves through and out the distal end **712** of the delivery catheter **710** to be positioned at the desired site adjacent the defect **104**.

After the rolled bladder **705** reaches the desired lateral position adjacent the defect **104**, the delivery catheter **710** may be removed from the patient's leg **101**. The rolled

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bladder **705** may then be unrolled in the opposite direction shown in FIG. **7B** to a relatively flattened shape with the cavity **130** positioned adjacent the defect **104**. When the first and second conduits **110**, **112** are reconnected to the reduced pressure source **115** and the first fluid supply **111**, a positive pressure is applied by the bracing fluid to the bracing chamber **107** so that it expands and exerts a force on the tibia **103** and on the interior wall **108** as described above while at the same time applying a reduced pressure to the reduced-pressure chamber **106** so that the opening **128** in the reduced-pressure chamber **106** (see FIG. **2B**) substantially seals the defect **104** as shown in FIG. **1A** and described in detail above. The positive pressure applied to the bracing chamber **107** and the weight of the patient being applied by the femur **102** through the manifold structure **120** and the interior wall **108** cause the exterior wall **127** of the bracing chamber **107** to mold to the curvature of the tibia **103** and the surrounding body tissues. Application of the positive pressure to the bracing chamber **107** also forces the interior wall **108** and the manifold structure **120** against the distal end of the femur to force the opening **128** of the reduced-pressure chamber **106** against the defect **104** after which a reduced pressure is applied to the defect **104** via the cavity **130**.

As described and shown in FIGS. **1A**, **4**, and **5** above, the cavity **130** may be filled with the scaffold component **422** or covered by a barrier material **520** closing the opening of the cavity **130**. Regardless of the structure, a reduced pressure is ultimately applied via the opening **128** in the reduced-pressure chamber **106** to the defect **104** so that the reduced pressure increases the seepage of blood and bone marrow through the fractures created by the microfracture surgery to draw blood and other stimulatory agents to the defect site. The reduced pressure therapy or treatment applied to the defect **104** depends on the size and shape of the defect **104** and various other factors known in the art. Consequently, the reduced pressure being applied may be continuous, variable at a specific frequency, or generally cyclical over time depending on the desired treatment for the patient and the characteristics of the defect **104**. The opening **128** in the reduced-pressure chamber **106** not only applies the reduced pressure directly to the defect **104**, but also mitigates the removal of the synovial fluid from the knee joint while the reduced pressure is being applied to the defect **104**.

Referring to FIG. **9**, another embodiment of a bladder **905** is shown that is substantially similar to the bladder **105** described above including the same components identified with similar reference numerals. The bladder **905** comprises a reduced-pressure chamber **906** and a bracing chamber **907** separated by an interior wall **908**, each chamber having an exterior wall **926** and **927**, respectively, all as described above with respect to the bladder **105**. The reduced-pressure chamber **906** also includes a manifold structure **920** that functions in the same fashion as the manifold structure **420** described above that fills most of the reduced-pressure chamber **906**. The reduced-pressure chamber **906** and the bracing chamber **907** each comprise an elongated conduit **914** and **916**, respectively, fluidly coupled to each chamber and extending from the bladder **905** through the skin of the leg **101** of the patient. The elongated conduits **914**, **916** may be an integrated portion of the reduced-pressure exterior wall **126** and the bracing exterior wall **127**, respectively, with the interior wall **908** extending therebetween. The manifold structure **920** extends through and substantially fills the elongated conduit **914** of the reduced-pressure chamber **906**. The distal ends of the elongated conduits **914**, **916** may be fluidly coupled to the first conduit **110** and the second conduit **112**, respectively, for providing a reduced

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pressure to the reduced-pressure chamber 906 and bracing fluid under a positive pressure to the bracing chamber 907 which function in the same manner as described above. The portion of the manifold structure 920 extending through the elongated conduit 914 of the reduced-pressure chamber 906 provides additional structural support for the reduced-pressure chamber 906 so that the bladder 905 can be more easily flattened or rolled as described above to facilitate insertion of the bladder 905 into the delivery catheter 710 (not shown).

Referring to FIG. 10A, a perspective view of the bladder 105 and the first and second conduits 110, 112 is shown. The bladder 105 further comprises an elongated tab 109 extending from the end of the bladder 105 opposite the side that is fluidly coupled to the first and second conduits 110, 112 (see also FIG. 1A). The elongated tab 109 is useful for guiding and manipulating the bladder 105 to the desired lateral position adjacent the defect 104 as described above. More specifically, the arthroscope may be inserted completely through the knee joint to create a path for the elongated tab 109 which would be inserted in advance of the bladder 105 through the delivery catheter 710 and ultimately extend out the other side of the patient's leg 101 as shown in FIG. 1A. The elongated tab 109 may be affixed to the exterior wall 126 of the reduced-pressure chamber 106 to facilitate positioning of the opening 128 in the reduced-pressure chamber 106 adjacent the defect 104. The elongated tab 109 may be affixed to the bladder 105 either prior to being inserted into the knee joint, or at any later point during the surgery to facilitate positioning of the bladder 105. For example, the elongated tab may be inserted from the opposite side of the knee joint through the incisional pathway and into the distal end 712 of the delivery catheter 710 where it can be affixed to or screwed into the bladder 105. The elongated tab 109 then can be used to pull the bladder 105 through the delivery catheter 710 instead of using the first and second conduits 110, 112 to push the bladder 105 to the desired position.

The bladder 105 may also include a sheath 119 covering the first and second conduits 110, 112 and affixed directly to the bladder 105. The sheath 119 may be constructed of a material for providing additional strength to the first and second conduits 110, 112 for pushing the bladder 105 into the desired position adjacent the defect 104 as described above. The sheath 119 and the elongated tab 109 may also be a single continuous piece of material as shown in FIG. 108 to further facilitate pushing and/or pulling the bladder 105 to the desired lateral position adjacent the defect 104 and positioning the opening 128 in the reduced-pressure chamber 106 so that it fully contacts the defect 104.

The design and materials of the bladder chamber, delivery tubes, and the like may be any contemplated in other embodiments of the application and are dependent upon numerous factors, including, for example, the location and size of the tissue site, the pressure of the bracing chamber to maintain the position of the opening 128 in the reduced-pressure chamber 106 against the defect 104. It is contemplated further that for all of the embodiments described herein, the number, size and type of bladder chambers may vary, depending upon the shape of the bone that contains the one or more tissue sites. As such, to better expand the bracing chambers and seal the one or more supportive foam structures to the bone comprising the tissue site, the bladder chamber may have two or more bracing chambers.

The application is also directed to a method of administering a reduced pressure therapy to a tissue site as described above, but more generally according to the following steps. The method comprises performing microfracture surgery on

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a bone at a tissue site and then delivering an embodiment of the reduced pressure delivery system described above to the tissue site. The method then comprises delivering a bracing substance to the bracing chamber such that the tissue site contact surface is in contact with the tissue site and delivering a reduced pressure to the reduced pressure chamber causing reduced pressure to be applied to the tissue site. This method can be used on any tissue site that is subjected to microfracture surgery. In some embodiments, the tissue site may be located in the knee joint as described above.

In additional embodiments, the application is directed to a method of performing microfracture surgery to a knee. The method comprises creating at least one microfracture in a bone at the base of an articular cartilage defect in the knee, then applying reduced pressure to the site of the defect. The reduced pressure in these embodiments can be applied by any means known in the art, and through any known system capable of delivering reduced pressure to the site. In some embodiments, the reduced pressure is applied using any appropriate embodiment of the reduced pressure delivery system described above.

As various changes could be made in any of the systems, apparatuses and methods described above without departing from the scope of the inventions, it is intended that all matter contained in the above description and shown in the accompanying drawings shall be interpreted as illustrative and not in a limiting sense.

All references cited in this specification are hereby incorporated by reference. The discussion of the references herein is intended merely to summarize the assertions made by the applicants and no admission is made that any reference constitutes prior art. Applicants reserve the right to challenge the accuracy and pertinence of the cited references.

We claim:

1. A system for applying a treatment to a defect in a first bone or a second bone of two bones forming a joint, the system comprising:

- a reduced-pressure source for providing a reduced pressure;
- a positive-pressure source for providing a positive pressure;
- a bladder formed of (i) a reduced-pressure chamber formed by flexible walls having an opening adapted to substantially surround the defect when positioned against the first bone and having a port in fluid communication with the reduced-pressure source; and (ii) a bracing chamber formed by flexible walls and adapted to be positioned against the second bone and having a port in fluid communication with the positive-pressure source for providing positive pressure to prevent the bracing chamber from collapsing when positive pressure is provided thereto; wherein a portion of the walls of each of the reduced-pressure chamber and the bracing chamber form an interior wall of the bladder; and
- a manifold disposed within the reduced-pressure chamber between the interior wall and the opening in the reduced-pressure chamber, and formed of a porous material adapted to distribute reduced pressure to the defect and provide structural support between the first bone and the interior wall to prevent the reduced-pressure chamber from collapsing when reduced pressure is provided thereto.

2. The system of claim 1, wherein the manifold partially fills the reduced-pressure chamber and includes a surface exposed through the opening to form a cavity adapted to be positioned adjacent the defect.

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3. The system of claim 2, wherein the manifold is a foam material in fluid communication with the defect.

4. The system of claim 3, wherein the foam material comprises a plurality of open cells having a size ranging from about 100 microns to about 1,500 microns.

5. The system of claim 2, wherein the manifold further comprises a scaffold component sized to fill the defect to facilitate new tissue formation.

6. The system of claim 2, wherein the interior wall is a common wall between the bracing chamber and the reduced-pressure chamber.

7. The system of claim 6, wherein the interior wall is substantially rigid.

8. The system of claim 2, further comprising a foam structure positioned in the cavity to provide additional bracing between the first bone and the bracing chamber.

9. The system of claim 8, wherein the foam structure is bioabsorbable.

10. The system of claim 8, wherein the foam structure serves as a scaffold for tissue growth and/or regeneration.

11. The system of claim 10, wherein the foam material is coated or infused with growth factors or nutrients.

12. The system of claim 8, wherein the foam structure adheres to the defect.

13. The system of claim 1, wherein the manifold substantially fills the reduced-pressure chamber and includes a surface exposed through the opening and adapted to be positioned adjacent the defect.

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14. The system of claim 13, wherein the manifold is a foam material in fluid communication with the defect.

15. The system of claim 14, wherein the foam material comprises a plurality of open cells having a size ranging from about 100 microns to about 1,500 microns.

16. The system of claim 13, wherein the interior wall is a common wall between the bracing chamber and the reduced-pressure chamber.

17. The system of claim 16, wherein the interior wall is substantially flexible.

18. The system of claim 13, wherein the manifold further comprises a scaffold component sized to fill the defect to facilitate tissue formation.

19. The system of claim 18, wherein the scaffold component provides additional bracing between the first bone and the bracing chamber.

20. The system of claim 18, wherein the scaffold component is bioabsorbable.

21. The system of claim 20, wherein the scaffold component adheres to the defect.

22. The system of claim 18, wherein the scaffold component is coated or infused with growth factors or nutrients.

23. The system of claim 1, wherein the walls of the reduced pressure chamber and the bracing chamber are comprised of one or more elastomers.

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